

10604022

| Ref # | Hits | Search Query   | DBs                                | Default Operator | Plurals | Time Stamp       |
|-------|------|--|------------------------------------|------------------|---------|------------------|
| L1    | 1    | deprotection near3 microwave   | US-PGPUB; USPAT; EPO; JPO; DERWENT | OR               | ON      | 2005/11/12 11:28 |
| L2    | 4    | deprotecting near3 microwave   | US-PGPUB; USPAT; EPO; JPO; DERWENT | OR               | ON      | 2005/11/12 11:29 |
| L3    | 794  | (deprotecting or deprotection or protect\$5) near3 microwave                 | US-PGPUB; USPAT; EPO; JPO; DERWENT | OR               | ON      | 2005/11/12 11:29 |
| L4    | 4    | (deprotecting or deprotection or de-protect\$4) near3 microwave              | US-PGPUB; USPAT; EPO; JPO; DERWENT | OR               | ON      | 2005/11/12 11:31 |
| L5    | 4    | ("peptide synthesis") near3 microwave  | US-PGPUB; USPAT; EPO; JPO; DERWENT | OR               | ON      | 2005/11/12 11:30 |
| L6    | 7    | (deprotecting or deprotection or de-protect\$4) near10 microwave             | US-PGPUB; USPAT; EPO; JPO; DERWENT | OR               | ON      | 2005/11/12 11:31 |
| L7    | 939  | (deprotecting or deprotection or de-protect\$4) and microwave                | US-PGPUB; USPAT; EPO; JPO; DERWENT | OR               | ON      | 2005/11/12 11:31 |
| L8    | 611  | (deprotecting or deprotection or de-protect\$4) and microwave and peptides   | US-PGPUB; USPAT; EPO; JPO; DERWENT | OR               | ON      | 2005/11/12 11:32 |
| L9    | 4    | (deprotecting or deprotection or de-protect\$4) and microwave near3 peptides | US-PGPUB; USPAT; EPO; JPO; DERWENT | OR               | ON      | 2005/11/12 11:33 |
| L10   | 0    | ((bmoc) near microwave)near3 peptides  | US-PGPUB; USPAT; EPO; JPO; DERWENT | OR               | ON      | 2005/11/12 11:33 |
| L11   | 4    | deprotect\$3 near3 microwave   | US-PGPUB; USPAT; EPO; JPO; DERWENT | OR               | ON      | 2005/11/12 12:29 |
| L12   | 3465 | coupling near3 microwave   | US-PGPUB; USPAT; EPO; JPO; DERWENT | OR               | ON      | 2005/11/12 12:30 |

|     |        |  |   |    |    |                  |
|-----|--------|--|---|----|----|------------------|
| L13 | 108216 | I12 and spp or ("solid phase")                   | US-PGPUB;<br>USPAT;<br>EPO; JPO;<br>DERWENT | OR | ON | 2005/11/12 12:30 |
| L14 | 37     | I12 and (spps or ("solid phase"))                | US-PGPUB;<br>USPAT;<br>EPO; JPO;<br>DERWENT | OR | ON | 2005/11/12 12:31 |
| L15 | 13     | I12 and (spps or ("solid phase"))<br>and peptide | US-PGPUB;<br>USPAT;<br>EPO; JPO;<br>DERWENT | OR | ON | 2005/11/12 12:31 |

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NEWS 15 OCT 27 EPFULL enhanced with additional content

NEWS EXPRESS JUNE 13 CURRENT WINDOWS VERSION IS V8.0, CURRENT MACINTOSH VERSION IS V6.0c(ENG) AND V6.0Jc(JP), AND CURRENT DISCOVER FILE IS DATED 13 JUNE 2005

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=> deprotection (3a) microwave  
L1 77 DEPROTECTION (3A) MICROWAVE

=> l1 and peptide  
L2 5 L1 AND PEPTIDE

=> dup rem  
ENTER L# LIST OR (END):12  
PROCESSING COMPLETED FOR L2  
L3 4 DUP REM L2 (1 DUPLICATE REMOVED)

=> d ibib abs total

L3 ANSWER 1 OF 4 CAPLUS COPYRIGHT 2005 ACS on STN  
ACCESSION NUMBER: 2005:560667 CAPLUS  
DOCUMENT NUMBER: 143:212166  
TITLE: MW-enhanced high-speed deprotection of Boc group using p-TsOH and concomitant formation of N-Me-amino acid benzyl ester p-TsOH salts  
AUTHOR(S): Babu, Vommina; Patil, Basanagoud; Vasanthakumar, Ganga-Ramu  
CORPORATE SOURCE: Department of Studies in Chemistry, Bangalore University, Bangalore, India  
SOURCE: Synthetic Communications (2005), 35(13), 1795-1802  
CODEN: SYNCV; ISSN: 0039-7911  
PUBLISHER: Taylor & Francis, Inc.  
DOCUMENT TYPE: Journal  
LANGUAGE: English

AB A high-speed, complete deprotection of Boc group from Boc (Boc = tert-butoxycarbonyl) amino acids and protected peptide esters employing p-TsOH in toluene under microwave irradiation is found to be complete in 30 s. The deprotection can be carried out in methanol and acetonitrile also. Under the present conditions, C-peptide benzyl esters and O-benzyl ethers have been found to be stable. This has permitted us to carry out the synthesis of [Leu]enkephalin employing the Boc/Bzl-group strategy. Further more, it has been found that both  $\text{Na}-\text{Fmoc}$  (Fmoc = 9-fluorenylmethoxycarbonyl) and  $\text{Na}-\text{Z}$  ( $\text{Z} =$  benzyloxycarbonyl) groups are completely stable. The present conditions can be extended for the concomitant removal of the Boc group and the formation of C-benzyl amino acid esters as well. This has been utilized for the synthesis of N-Me amino acid benzyl esters starting from Boc-N-Me amino acids in a single step.

REFERENCE COUNT: 29 THERE ARE 29 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L3 ANSWER 2 OF 4 CAPLUS COPYRIGHT 2005 ACS on STN  
ACCESSION NUMBER: 2004:658783 CAPLUS  
TITLE: Microwave-enhanced solid-phase peptide synthesis  
AUTHOR(S): Collins, Jonathan M.  
CORPORATE SOURCE: CEM Corporation, Matthews, NC, 28106-0200, USA  
SOURCE: Abstracts of Papers, 228th ACS National Meeting, Philadelphia, PA, United States, August 22-26, 2004

(2004), ORGN-518. American Chemical Society:  
Washington, D. C.  
CODEN: 69FTZ8

DOCUMENT TYPE: Conference; Meeting Abstract  
LANGUAGE: English

AB Microwave energy has proven to be a valuable tool for organic synthesis. Recently, microwave has been used for enhanced Fmoc solid phase peptide synthesis. With **microwave** energy, **deprotection** and coupling reactions can be performed in 3 and 4 min resp. This paper builds on previous work and demonstrates the successful application of microwave energy for longer 30-40 amino acid peptide sequences. Variation in deprotection and coupling chemistries will be presented and discussed.

L3 ANSWER 3 OF 4 EMBASE COPYRIGHT (c) 2005 Elsevier B.V. All rights reserved on STN DUPLICATE 1

ACCESSION NUMBER: 2001246403 EMBASE  
TITLE: Rapid **microwave-assisted deprotection** of N-Cbz and N-Bn derivatives.  
AUTHOR: Daga M.C.; Taddei M.; Varchi G.  
CORPORATE SOURCE: M. Taddei, Dipartimento di Chimica, Universita degli Studi di Sassari, Via Vienna 2, 07100 Sassari, Italy  
SOURCE: Tetrahedron Letters, (30 Jul 2001) Vol. 42, No. 31, pp. 5191-5194.  
Refs: 15  
ISSN: 0040-4039 CODEN: TELEAY  
PUBLISHER IDENT.: S 0040-4039(01)00969-8  
COUNTRY: United Kingdom  
DOCUMENT TYPE: Journal; Article  
FILE SEGMENT: 029 Clinical Biochemistry  
LANGUAGE: English  
SUMMARY LANGUAGE: English  
ENTRY DATE: Entered STN: 20010802  
Last Updated on STN: 20010802

AB Catalytic-transfer hydrogenation in iso-propanol under microwave irradiation has been performed to rapidly deprotect N-Cbz and N-Bn derivatives. The method is particularly suitable for the synthesis of short **peptides** and can also be carried out on supported molecules. The rapid cleavage of chiral molecules derived from (S)-1-phenylethylamine can be very useful for asymmetric synthesis of nitrogen containing compounds. .COPYRGT. 2001 Elsevier Science Ltd.

L3 ANSWER 4 OF 4 CAPLUS COPYRIGHT 2005 ACS on STN

ACCESSION NUMBER: 2001:494797 CAPLUS  
DOCUMENT NUMBER: 135:242481  
TITLE: Utilization of microwave technique for cleavage of acid-labile groups in **peptide** chemistry  
AUTHOR(S): Sebestik, Jaroslav; Hlavacek, Jan; Stibor, Ivan  
CORPORATE SOURCE: Ustav Org. Chem. a Biochem., Akad. Ved Ceske Republiky, Prague, 166 10, Czech Rep.  
SOURCE: Chemicke Listy (2001), 95(6), 365-367  
PUBLISHER: Ceska Spolecnost Chemicka  
DOCUMENT TYPE: Journal  
LANGUAGE: Czech

AB We report our findings on the microwave-assisted silica gel removal of Boc and t-Bu groups protecting  $\alpha$ -NH<sub>2</sub> and COOH groups in amino acids and **peptides**, and also on selectivity of this deprotection to OH-protecting t-Bu groups. To obtain high conversions in **deprotection**, increased **microwave** doses and prolongation of the reaction time are required. As a consequence, relatively large amts. of side products were detected. The microwave radiation could possibly be utilized in deprotection of **peptides** when the trifluoroacetic acid cleavage and subsequent neutralization of the acid with a tertiary base have to be avoided.

=> d his

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FILE 'MEDLINE, BIOSIS, EMBASE, CAPLUS' ENTERED AT 11:36:01 ON 12 NOV 2005  
L1 77 DEPROTECTION (3A) MICROWAVE  
L2 5 L1 AND PEPTIDE  
L3 4 DUP REM L2 (1 DUPLICATE REMOVED)

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L1 77 DEPROTECTION (3A) MICROWAVE  
L2 5 L1 AND PEPTIDE  
L3 4 DUP REM L2 (1 DUPLICATE REMOVED)

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| FULL ESTIMATED COST  | 19.47            | 19.68         |

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=> l1 and "solid phase"  
L4 2 L1 AND "SOLID PHASE"

=> dup rem 14  
PROCESSING COMPLETED FOR L4  
L5 2 DUP REM L4 (0 DUPLICATES REMOVED)

=> d ibib abs total

L5 ANSWER 1 OF 2 CAPLUS COPYRIGHT 2005 ACS on STN  
ACCESSION NUMBER: 2004:658783 CAPLUS  
TITLE: Microwave-enhanced **solid-phase**  
peptide synthesis  
AUTHOR(S): Collins, Jonathan M.  
CORPORATE SOURCE: CEM Corporation, Matthews, NC, 28106-0200, USA  
SOURCE: Abstracts of Papers, 228th ACS National Meeting,  
Philadelphia, PA, United States, August 22-26, 2004  
(2004), ORGN-518. American Chemical Society:  
Washington, D. C.  
CODEN: 69FTZ8

DOCUMENT TYPE: Conference; Meeting Abstract

LANGUAGE: English

AB Microwave energy has proven to be a valuable tool for organic synthesis.  
Recently, microwave has been used for enhanced Fmoc **solid**  
**phase** peptide synthesis. With **microwave** energy,  
**deprotection** and coupling reactions can be performed in 3 and 4  
min resp. This paper builds on previous work and demonstrates the  
successful application of microwave energy for longer 30-40 amino acid  
peptide sequences. Variation in deprotection and coupling chemistries  
will be presented and discussed.

L5 ANSWER 2 OF 2 BIOSIS COPYRIGHT (c) 2005 The Thomson Corporation on STN  
ACCESSION NUMBER: 1998:490462 BIOSIS  
DOCUMENT NUMBER: PREV199800490462  
TITLE: Cleavage of oligodeoxyribonucleotides from polymer supports  
and their rapid **deprotection** under  
**microwaves**.  
AUTHOR(S): Gupta, K. C. [Reprint author]; Kumar, P.  
CORPORATE SOURCE: Nucleic Acids Res. Lab., Centre Biochem. Technol., Mall  
Rd., Delhi Univ. Campus, Delhi 110 007, India  
SOURCE: Nucleosides and Nucleotides, (Sept.-Nov., 1998) Vol. 17,  
No. 9-11, pp. 1761-1766. print.  
CODEN: NUNUD5. ISSN: 0732-8311.  
DOCUMENT TYPE: Article  
LANGUAGE: English  
ENTRY DATE: Entered STN: 18 Nov 1998  
Last Updated on STN: 18 Nov 1998

AB Novel conditions for the cleavage of oligodeoxynucleotides from polymer supports and their complete **deprotection** under **microwaves** have been developed. The oligonucleotides synthesized using phosphoramidite synthons carrying base labile (Pac, Dmf and t-Bpac) and conventional (Bz for A and C and Pac for G) protecting groups for nucleic bases were deprotected using 0.2M sodium hydroxide (MeOH:H<sub>2</sub>O::1:1, v/v) = Reagent A and 1M sodium hydroxide (MeOH:H<sub>2</sub>O::1:1, v/v) = Reagent B, respectively under microwaves. The deprotected oligonucleotides were found to be comparable with the corresponding oligonucleotides deprotected under standard conditions (aqueous ammonia at 55degreeC).

=> activation (3a) microwave  
L6 530 ACTIVATION (3A) MICROWAVE

=> 16 and peptide and "solid phase"  
L7 1 L6 AND PEPTIDE AND "SOLID PHASE"

=> d ibib abs total

L7 ANSWER 1 OF 1 CAPLUS COPYRIGHT 2005 ACS on STN  
ACCESSION NUMBER: 2003:1001899 CAPLUS  
DOCUMENT NUMBER: 140:236083  
TITLE: Synthesis of methyleneaminodipeptides via ring opening of a 2-(t-butoxycarbonylmethyl)aziridine derivative  
AUTHOR(S): Thierry, Josiane; Servajean, Vincent  
CORPORATE SOURCE: Institut de Chimie des Substances Naturelles, CNRS, Gif-sur-Yvette, 91198, Fr.  
SOURCE: Tetrahedron Letters (2004), 45(4), 821-823  
CODEN: TELEAY; ISSN: 0040-4039  
PUBLISHER: Elsevier Science B.V.  
DOCUMENT TYPE: Journal  
LANGUAGE: English  
OTHER SOURCE(S): CASREACT 140:236083

AB The reactivity of 2-(tert-butoxycarbonylmethyl)aziridine-1-carboxylic acid benzyl ester has been studied with various N-nucleophiles. The ring-opening reaction was always regioselective, the nucleophile attacking preferentially the less hindered carbon of the aziridine. The reaction was used to prepare a methyleneamino pseudodipeptide using the  $\alpha$ -amine of a lysine ester. The solvent-free reaction of 2-(tert-butoxycarbonylmethyl)aziridine derivative with benzylamine under **microwave activation** on solid support gave the same result as the classical reaction but in a much shorter time and represents a significant improvement in the procedure.

REFERENCE COUNT: 12 THERE ARE 12 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

=> activation and protection  
L8 37668 ACTIVATION AND PROTECTION

=> 18 and microwave and peptide  
L9 0 L8 AND MICROWAVE AND PEPTIDE

=> s daga, maria caterina/au  
L10 0 DAGA, MARIA CATERINA/AU

=> daga, caterina/au  
L11 1 DAGA, CATERINA/AU

=> taddei, maurizio/au  
L12 162 TADDEI, MAURIZIO/AU

=> varchi, greta/au  
L13 23 VARCHI, GRETA/AU

. => l12 and l13  
L14 0 L12 AND L13

=> l12 and "solid phase"  
L15 31 L12 AND "SOLID PHASE"

=> l15 and microwave  
L16 5 L15 AND MICROWAVE

=> dup rem l16  
PROCESSING COMPLETED FOR L16  
L17 3 DUP REM L16 (2 DUPLICATES REMOVED)

=> d ibib abs total

L17 ANSWER 1 OF 3 MEDLINE on STN DUPLICATE 1  
ACCESSION NUMBER: 2003450794 MEDLINE  
DOCUMENT NUMBER: PubMed ID: 14510574  
TITLE: Solid-phase synthesis of conformationally constrained peptidomimetics based on a 3,6-disubstituted-1,4-diazepan-2,5-dione core.  
AUTHOR: Lampariello Lucia Raffaella; Piras Daniela; Rodriguez Manuela; Taddei Maurizio  
CORPORATE SOURCE: Dipartimento di Chimica, Universita degli Studi di Sassari, Via Vienna 2, I-07100 Sassari, Italy.  
SOURCE: Journal of organic chemistry, (2003 Oct 3) 68 (20) 7893-5.  
Journal code: 2985193R. ISSN: 0022-3263.  
PUB. COUNTRY: United States  
DOCUMENT TYPE: Journal; Article; (JOURNAL ARTICLE)  
LANGUAGE: English  
FILE SEGMENT: Priority Journals  
ENTRY MONTH: 200404  
ENTRY DATE: Entered STN: 20030928  
Last Updated on STN: 20040409  
Entered Medline: 20040408  
AB Starting from a Cl-trytyl linked hydroxylamine, a hydroxamic dipeptide having serine in the second position was prepared by using DMTMM as the coupling agent. Mitsunobu cyclization carried out under **microwave** heating gave very good yields of a 3,6-disubstituted-perhydro-diazepin-2,5-dione. This heterocycle can be used as a new platform for combinatorial chemistry or as a constraint to rigidify a small peptide.

L17 ANSWER 2 OF 3 CAPLUS COPYRIGHT 2005 ACS on STN  
ACCESSION NUMBER: 2003:190685 CAPLUS  
DOCUMENT NUMBER: 139:53283  
TITLE: A new, rapid, general procedure for the synthesis of organic molecules supported on methoxy-polyethylene glycol (MeOPEG) under **microwave** irradiation conditions  
AUTHOR(S): Porcheddu, Andrea; Ruda, Gian Filippo; Sega, Alessandro; Taddei, Maurizio  
CORPORATE SOURCE: Dipartimento di Chimica, Universita degli Studi di Sassari, Sassari, 07100, Italy  
SOURCE: European Journal of Organic Chemistry (2003), (5), 907-912  
PUBLISHER: Wiley-VCH Verlag GmbH & Co. KGaA  
DOCUMENT TYPE: Journal  
LANGUAGE: English  
OTHER SOURCE(S): CASREACT 139:53283  
AB The procedure for the precipitation of mols. supported on MeOPEG (mol. mass 5000) and their purification by fractional crystallization has been made easier by use of

microwave irradiation A correct choice of the solvent employed for reaction or purification (DME, THF, 1,2-dichlorobenzene, iPrOH, ethylene glycol) allows working with 10 g of MeOPEG-OH, dissolved in 100 mL of solvent, under microwave irradiation conditions and for crystallization to be induced just by removal of the reaction flask from the microwave oven. No addnl. precipitation solvents are needed, thus reducing the reaction times and the potential hazards of working with large amts. of flammable solvents. The syntheses of several peptides and of a tetrasubstituted pyridine are reported. Large amts. of MeOPEG-OH may be used in this procedure, and so polyethylene glycol assisted organic synthesis can be regarded as a valid preparative technique.

REFERENCE COUNT: 46 THERE ARE 46 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L17 ANSWER 3 OF 3 CAPLUS COPYRIGHT 2005 ACS on STN

ACCESSION NUMBER: 2003:444197 CAPLUS

DOCUMENT NUMBER: 139:164736

TITLE: Cellulose Beads: a New Versatile Solid Support for Microwave-Assisted Synthesis. Preparation of Pyrazole and Isoxazole Libraries

AUTHOR(S): De Luca, Lidia; Giacomelli, Giampaolo; Porcheddu, Andrea; Salaris, Margherita; Taddei, Maurizio

CORPORATE SOURCE: Dipartimento di Chimica, Universita degli Studi di Sassari, Sassari, I-07100, Italy

SOURCE: Journal of Combinatorial Chemistry (2003), 5(4), 465-471

CODEN: JCCHFF; ISSN: 1520-4766

PUBLISHER: American Chemical Society

DOCUMENT TYPE: Journal

LANGUAGE: English

OTHER SOURCE(S): CASREACT 139:164736

AB Combinatorial libraries of pyrazoles and isoxazoles are prepared by cyclocondensation reactions of  $\beta$ -ketoesters and  $\beta$ -ketoamides, 1-(dimethoxymethyl)imidazole, and hydrazines or hydroxylamine on solid-phase using a novel aminophenyl-substituted cellulose resin. Heating the aminophenyl-substituted cellulose with either  $\beta$ -ketoesters or  $\beta$ -ketoamides and 1-(dimethoxymethyl)imidazole yields polymer-bound enaminones in >99% yields by colorimetric assays; heating the resin-bound enaminones with hydrazines or hydroxylamine in isopropanol yields the product heterocycles in addition to the aminophenyl-substituted cellulose resin which can be reused. Testing of the resin with  $\beta$ -naphthol and sodium nitrite gives a red color if free arylamino groups are present on the resin, while testing with iron (III) chloride allows the presence of resin-bound  $\beta$ -enaminone moieties to be determined. The added stability of cellulose to thermal shock allows both conventional and microwave heating to be used for solid-phase reactions. One-pot and multiple step reactions are used to obtain pyrazoles and isoxazoles in 97-99% yields; one-pot synthesis using microwave irradiation gives the heterocyclic products in >95% yields and in >98% purities.

REFERENCE COUNT: 105 THERE ARE 105 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

=> ("solid phase" (3a) peptides) or SPPS

L18 4249 ("SOLID PHASE" (3A) PEPTIDES) OR SPPS

=> (activation or deprotection)

L19 2167208 (ACTIVATION OR DEPROTECTION)

=> 118 and 119

L20 535 L18 AND L19

=> 120 and microwave

L21 1 L20 AND MICROWAVE

=> d ibib abs total

L21 ANSWER 1 OF 1 CAPLUS COPYRIGHT 2005 ACS on STN  
ACCESSION NUMBER: 2004:227419 CAPLUS  
TITLE: Effect of **microwave** energy on solid phase peptide synthesis  
AUTHOR(S): Collins, Jonathan M.; Hassman, C. Fred; King, Edward E.; Lambert, Joseph  
CORPORATE SOURCE: Life Sciences Division, CEM Corporation, Matthews, NC, 28106, USA  
SOURCE: Abstracts of Papers, 227th ACS National Meeting, Anaheim, CA, United States, March 28-April 1, 2004 (2004), ORGN-549. American Chemical Society: Washington, D. C.  
CODEN: 69FGKM  
DOCUMENT TYPE: Conference; Meeting Abstract  
LANGUAGE: English  
AB The application of **microwave** energy has proved to be a major enabling tool for many chemical applications requiring energy input. A new automated system for **microwave** assisted solid phase peptide synthesis has been developed that allows for complete cycle times of ten minutes as well as final peptide cleavage in ten minutes. A single mode cavity is used to allow for a high **microwave** power d. and a uniform field distribution. The stability of activated amino acids under **microwave** irradiation was investigated using PyBOP and HBTU activation. The effect of **microwave** energy on conventional side reactions with SPPS such as racemization and aspartimide formation was investigated and found to compare very favorably with conventional methods. Also, exciting changes in coupling chemistries possible with **microwave** energy will be presented that help to further suppress racemization. The application of this new method will be shown on a variety of peptide sequences.

=> (pybop or hbtu or hatu or pyaop or hobt)  
L22 2370 (PYBOP OR HBTU OR HATU OR PYAOP OR HOBT)

=> 122 and 118 and microwave  
L23 1 L22 AND L18 AND MICROWAVE

=> d

L23 ANSWER 1 OF 1 CAPLUS COPYRIGHT 2005 ACS on STN  
AN 2004:227419 CAPLUS  
TI Effect of **microwave** energy on solid phase peptide synthesis  
AU Collins, Jonathan M.; Hassman, C. Fred; King, Edward E.; Lambert, Joseph  
CS Life Sciences Division, CEM Corporation, Matthews, NC, 28106, USA  
SO Abstracts of Papers, 227th ACS National Meeting, Anaheim, CA, United States, March 28-April 1, 2004 (2004), ORGN-549 Publisher: American Chemical Society, Washington, D. C.  
CODEN: 69FGKM  
DT Conference; Meeting Abstract  
LA English

=> (pybop or hbtu or hatu or pyaop or hobt) and microwave  
L24 9 (PYBOP OR HBTU OR HATU OR PYAOP OR HOBT) AND MICROWAVE

=> dup rem 124  
PROCESSING COMPLETED FOR L24  
L25 6 DUP REM L24 (3 DUPLICATES REMOVED)

=> d ibib abs total

L25 ANSWER 1 OF 6 CAPLUS COPYRIGHT 2005 ACS on STN  
ACCESSION NUMBER: 2005:702713 CAPLUS  
TITLE: **Microwave** accelerated high speed solution synthesis of peptides employing **HATU/HOAt**  
AUTHOR(S): Sudarshan, Naremaddepalli S.; Babu, Vommina V. Suresh  
CORPORATE SOURCE: Department of Studies in Chemistry Central College Campus, Bangalore University, Bangalore, 560 001, India  
SOURCE: Indian Journal of Chemistry, Section B: Organic Chemistry Including Medicinal Chemistry (2005), 44B(7), 1509-1511  
CODEN: IJSBDB; ISSN: 0376-4699  
PUBLISHER: National Institute of Science Communication and Information Resources  
DOCUMENT TYPE: Journal  
LANGUAGE: English  
AB The chemical synthesis of peptides employing **HATU/HOAt** as a coupling agent under **microwave** irradiation has been described. The coupling is found to be complete in 30 - 40 s. The yield as well as purity of the peptides made is found to be good. All the peptides prepared have been characterized by <sup>1</sup>H NMR and mass spectroscopy.  
REFERENCE COUNT: 21 THERE ARE 21 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L25 ANSWER 2 OF 6 EMBASE COPYRIGHT (c) 2005 Elsevier B.V. All rights reserved on STN DUPLICATE 1  
ACCESSION NUMBER: 2005104988 EMBASE  
TITLE: A reactivity test for **HBTU**-activated carboxylic acids with low reactivity and competitive coupling of N-methylpyrrole derivatives.  
AUTHOR: Ernst T.; Richert C.  
CORPORATE SOURCE: C. Richert, Institute for Organic Chemistry, University of Karlsruhe (TH), 76131 Karlsruhe, Germany. cr@rrg.uka.de  
SOURCE: Synlett, (16 Feb 2005) No. 3, pp. 411-416.  
Refs: 36  
ISSN: 0936-5214 CODEN: SYNLES  
COUNTRY: Germany  
DOCUMENT TYPE: Journal; Article  
FILE SEGMENT: 029 Clinical Biochemistry  
LANGUAGE: English  
SUMMARY LANGUAGE: English  
ENTRY DATE: Entered STN: 20050324  
Last Updated on STN: 20050324  
AB N-Methylpyrrole carboxylic acids are building blocks for oligopyrroleamides that bind DNA duplexes via the minor groove. The reactivity of **HBTU**-based active esters of four methylpyrroles in amide-forming reactions was determined. When assayed against **HBTU**-activated N-acetylleucine, a 6-250-fold lower reactivity was found. When assayed against the NHS ester of Boc-valine, the reactivity was up to 4-fold lower. Despite large differences in reactivity, mixed couplings were successfully performed with all four pyrroles, generating small libraries of modified oligonucleotides suitable for spectrometrically monitored selection experiments. **Microwave** irradiation accelerated coupling of an Fmoc-protected pyrrole to an amine on solid support. .COPYRGHT. Georg Thieme Verlag Stuttgart.

L25 ANSWER 3 OF 6 CAPLUS COPYRIGHT 2005 ACS on STN  
ACCESSION NUMBER: 2005:739910 CAPLUS  
TITLE: Chemoselective synthesis of 1,2,4-triazole derivatives using solid-supported reagents as selective inhibitors of cyclin-dependent kinase 4 (Cdk4)  
AUTHOR(S): Kim, Kyungjin; Chen, Li; Chen, Yingsi; Depinto, Wanda; Lovey, Allen; McComas, Warren; Xiang, Qing; Yin, Xuefeng

CORPORATE SOURCE: Discovery Chemistry, Hoffmann-La Roche, Inc, Nutley, NJ, 07110-1199, USA  
SOURCE: Abstracts of Papers, 230th ACS National Meeting, Washington, DC, United States, Aug. 28-Sept. 1, 2005 (2005), MEDI-402. American Chemical Society: Washington, D. C.  
CODEN: 69HFCL  
DOCUMENT TYPE: Conference; Meeting Abstract; (computer optical disk)  
LANGUAGE: English  
AB A series of 1,2,4-triazaole derivs. was identified as selective inhibitors of cyclin-dependent kinase 4 (CDK4). A novel synthetic pathway using a combination of solid-supported reagents and microwave-assisted technol. was explored to facilitate SAR development. Polymer-supported activated ester acylation reagents were prepared from a variety of com. available aromatic carboxylic acids with polymer-supported HOBt. Utilizing microwave technol. to accelerate reaction rates, N-acylation reactions were successfully explored. A simple purification method to isolate the N1 from N2 regioisomer of the 1,2,4-triazole scaffold was developed. Herein we will also discuss some analogs with low nanomolar activity toward CDK4 as well as greater than 10-fold selectivity against CDK1 and CDK2.

L25 ANSWER 4 OF 6 CAPLUS COPYRIGHT 2005 ACS on STN  
ACCESSION NUMBER: 2004:227419 CAPLUS  
TITLE: Effect of microwave energy on solid phase peptide synthesis  
AUTHOR(S): Collins, Jonathan M.; Hassman, C. Fred; King, Edward E.; Lambert, Joseph  
CORPORATE SOURCE: Life Sciences Division, CEM Corporation, Matthews, NC, 28106, USA  
SOURCE: Abstracts of Papers, 227th ACS National Meeting, Anaheim, CA, United States, March 28-April 1, 2004 (2004), ORGN-549. American Chemical Society: Washington, D. C.  
CODEN: 69FGKM  
DOCUMENT TYPE: Conference; Meeting Abstract  
LANGUAGE: English  
AB The application of microwave energy has proved to be a major enabling tool for many chemical applications requiring energy input. A new automated system for microwave assisted solid phase peptide synthesis has been developed that allows for complete cycle times of ten minutes as well as final peptide cleavage in ten minutes. A single mode cavity is used to allow for a high microwave power d. and a uniform field distribution. The stability of activated amino acids under microwave irradiation was investigated using PyBOP and HBTU activation. The effect of microwave energy on conventional side reactions with SPPS such as racemization and aspartimide formation was investigated and found to compare very favorably with conventional methods. Also, exciting changes in coupling chemistries possible with microwave energy will be presented that help to further suppress racemization. The application of this new method will be shown on a variety of peptide sequences.

L25 ANSWER 5 OF 6 CAPLUS COPYRIGHT 2005 ACS on STN  
ACCESSION NUMBER: 2003:552687 CAPLUS  
DOCUMENT NUMBER: 140:59480  
TITLE: Microwave-assisted coupling of carboxylic acids to a polymer bound hydrazine linker  
AUTHOR(S): Lindquist, Charlotta; Tedebark, Ulf; Ersoy, Oguz; Somfai, Peter  
CORPORATE SOURCE: Organic Chemistry, Department of Chemistry, Royal Institute of Technology, Stockholm, Swed.  
SOURCE: Synthetic Communications (2003), 33(13), 2257-2262  
PUBLISHER: Marcel Dekker, Inc.

DOCUMENT TYPE: Journal  
LANGUAGE: English  
OTHER SOURCE(S): CASREACT 140:59480  
AB A set of carboxylic acids, all being potential scaffolds for combinatorial chemical or peptide synthesis, were coupled to a polymer bound aryl hydrazine linker using **microwave** irradiation in good yields. Improved yields and reduced reaction times were achieved by using **microwave**-assisted heating compared to conventional heating.  
REFERENCE COUNT: 12 THERE ARE 12 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L25 ANSWER 6 OF 6 BIOSIS COPYRIGHT (c) 2005 The Thomson Corporation on STN  
DUPLICATE 2

ACCESSION NUMBER: 2001:427078 BIOSIS

DOCUMENT NUMBER: PREV200100427078

TITLE: **Microwave**-enhanced solution coupling of the alpha,alpha-dialkyl amino acid, Aib.

AUTHOR(S): Santagada, Vincenzo [Reprint author]; Fiorino, Ferdinando; Perissutti, Elisa; Severino, Beatrice; De Filippis, Vincenzo; Vivenzio, Beniamino; Caliendo, Giuseppe

CORPORATE SOURCE: Dipartimento di Chimica Farmaceutica e Tossicologica, Federico II, Universita di Napoli, Via D. Montesano, 49, 80131, Naples, Italy

SOURCE: *Tetrahedron Letters*, (30 July, 2001) Vol. 42, No. 31, pp. 5171-5173. print.

CODEN: TELEAY. ISSN: 0040-4039.

DOCUMENT TYPE: Article

LANGUAGE: English

ENTRY DATE: Entered STN: 12 Sep 2001

Last Updated on STN: 22 Feb 2002

AB The difficult coupling of alpha-aminoisobutyric acid (Aib), during the synthesis of dipeptides (1-6), was carried out using **PyBOP/HOBt** and **HBTU/HOBt** reagents by application of **microwave** energy in the presence of solvent. Room temperature, conventional heating (oil bath) and **microwave** irradiation of the reactions are compared. Synthesis by **microwave** irradiation gave the desired compounds in higher yields and in shorter reaction times than those obtained by conventional heating or at room temperature.

=> d his

(FILE 'HOME' ENTERED AT 11:35:43 ON 12 NOV 2005)

FILE 'MEDLINE, BIOSIS, EMBASE, CAPLUS' ENTERED AT 11:36:01 ON 12 NOV 2005

L1 77 DEPROTECTION (3A) MICROWAVE

L2 5 L1 AND PEPTIDE

L3 4 DUP REM L2 (1 DUPLICATE REMOVED)

FILE 'MEDLINE, BIOSIS, EMBASE, CAPLUS' ENTERED AT 11:51:37 ON 12 NOV 2005

L4 2 L1 AND "SOLID PHASE"

L5 2 DUP REM L4 (0 DUPLICATES REMOVED)

L6 530 ACTIVATION (3A) MICROWAVE

L7 1 L6 AND PEPTIDE AND "SOLID PHASE"

L8 37668 ACTIVATION AND PROTECTION

L9 0 L8 AND MICROWAVE AND PEPTIDE

L10 0 S DAGA, MARIA CATERINA/AU

L11 1 DAGA, CATERINA/AU

L12 162 TADDEI, MAURIZIO/AU

L13 23 VARCHI, GRETA/AU

L14 0 L12 AND L13

L15 31 L12 AND "SOLID PHASE"

L16 5 L15 AND MICROWAVE

L17 3 DUP REM L16 (2 DUPLICATES REMOVED)

L18 4249 ("SOLID PHASE" (3A) PEPTIDES) OR SPPS  
L19 2167208 (ACTIVATION OR DEPROTECTION)  
L20 535 L18 AND L19  
L21 1 L20 AND MICROWAVE  
L22 2370 (PYBOP OR HBTU OR HATU OR PYAOP OR HOBT)  
L23 1 L22 AND L18 AND MICROWAVE  
L24 9 (PYBOP OR HBTU OR HATU OR PYAOP OR HOBT) AND MICROWAVE  
L25 6 DUP REM L24 (3 DUPLICATES REMOVED)

=> 112 and polymeric or polymer or resin  
L26 2128248 L12 AND POLYMERIC OR POLYMER OR RESIN

=> 112 and (polymeric or polymer or resin)  
L27 28 L12 AND (POLYMERIC OR POLYMER OR RESIN)

=> 127 and microwave  
L28 7 L27 AND MICROWAVE

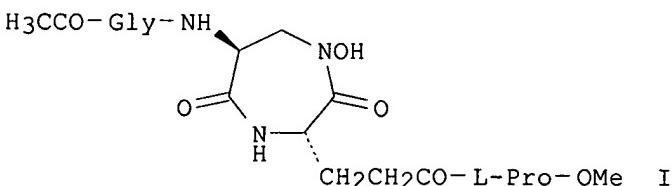
=> 127 and peptide  
L29 11 L27 AND PEPTIDE

=> 128 and peptide  
L30 2 L28 AND PEPTIDE

=> dup rem 130  
PROCESSING COMPLETED FOR L30  
L31 2 DUP REM L30 (0 DUPLICATES REMOVED)

=> d ibib abs total

L31 ANSWER 1 OF 2 CAPLUS COPYRIGHT 2005 ACS on STN  
ACCESSION NUMBER: 2003:679456 CAPLUS  
DOCUMENT NUMBER: 139:323781  
TITLE: Solid-Phase Synthesis of Conformationally Constrained Peptidomimetics Based on a 3,6-Disubstituted-1,4-diazepan-2,5-dione Core  
AUTHOR(S): Lampariello, Lucia Raffaella; Piras, Daniela; Rodriguez, Manuela; Taddei, Maurizio  
CORPORATE SOURCE: Dipartimento di Chimica, Universita degli Studi di Sassari, Sassari, I-07100, Italy  
SOURCE: Journal of Organic Chemistry (2003), 68(20), 7893-7895  
CODEN: JOCEAH; ISSN: 0022-3263  
PUBLISHER: American Chemical Society  
DOCUMENT TYPE: Journal  
LANGUAGE: English  
OTHER SOURCE(S): CASREACT 139:323781  
GI



AB Starting from a chlorotriptyl resin-linked hydroxylamine, a hydroxamic dipeptide having serine at the N-terminus was prepared by using DMTMM [4-(4,6-dimethoxy-1,3,5-triazin-2-yl)-4-methylmorpholinium chloride] as the coupling agent. Under **microwave** heating, Mitsunobu cyclization of the hydroxamic dipeptide gave a 3,6-disubstituted-perhydro-diazepin-2,5-dione in very good yields. Thus, by using

Fmoc-Ser-Glu(OCH<sub>2</sub>CH:CH<sub>2</sub>)-NH-O-**Resin**, H<sub>3</sub>CCO-Gly-OH and H-Pro-OMe,  
peptidomimetic I was prepared in four steps in 75% yield.

REFERENCE COUNT: 42 THERE ARE 42 CITED REFERENCES AVAILABLE FOR THIS  
RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L31 ANSWER 2 OF 2 CAPLUS COPYRIGHT 2005 ACS on STN  
ACCESSION NUMBER: 2003:190685 CAPLUS  
DOCUMENT NUMBER: 139:53283  
TITLE: A new, rapid, general procedure for the synthesis of  
organic molecules supported on methoxy-polyethylene  
glycol (MeOPEG) under **microwave** irradiation  
conditions  
AUTHOR(S): Porcheddu, Andrea; Ruda, Gian Filippo; Sega,  
Alessandro; Taddei, Maurizio  
CORPORATE SOURCE: Dipartimento di Chimica, Universita degli Studi di  
Sassari, Sassari, 07100, Italy  
SOURCE: European Journal of Organic Chemistry (2003), (5),  
907-912  
CODEN: EJOCFK; ISSN: 1434-193X  
PUBLISHER: Wiley-VCH Verlag GmbH & Co. KGaA  
DOCUMENT TYPE: Journal  
LANGUAGE: English  
OTHER SOURCE(S): CASREACT 139:53283  
AB The procedure for the precipitation of mols. supported on MeOPEG (mol. mass  
5000)  
and their purification by fractional crystallization has been made easier by  
use of  
**microwave** irradiation A correct choice of the solvent employed for  
reaction or purification (DME, THF, 1,2-dichlorobenzene, iPrOH, ethylene  
glycol) allows working with 10 g of MeOPEG-OH, dissolved in 100 mL of  
solvent, under **microwave** irradiation conditions and for crystallization to  
be induced just by removal of the reaction flask from the  
**microwave** oven. No addnl. precipitation solvents are needed, thus  
reducing the reaction times and the potential hazards of working with  
large amts. of flammable solvents. The syntheses of several  
**peptides** and of a tetrasubstituted pyridine are reported. Large  
amts. of MeOPEG-OH may be used in this procedure, and so polyethylene  
glycol assisted organic synthesis can be regarded as a valid preparative  
technique.  
REFERENCE COUNT: 46 THERE ARE 46 CITED REFERENCES AVAILABLE FOR THIS  
RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

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| ENTRY      | SESSION |

FULL ESTIMATED COST

151.11 170.79

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|------------|---------|
| ENTRY      | SESSION |

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|------------|---------|
| ENTRY      | SESSION |

FULL ESTIMATED COST

0.24 171.03

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|----------------------|------------------|---------------|
| COST IN U.S. DOLLARS | SINCE FILE ENTRY | TOTAL SESSION |
| FULL ESTIMATED COST  | 0.24             | 171.03        |

|  |                  |               |
|--|------------------|---------------|
| DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS) | SINCE FILE ENTRY | TOTAL SESSION |
| CA SUBSCRIBER PRICE                        | 0.00             | -10.22        |

=> coupling (3a) microwave  
    3 COUPLING  
    0 MICROWAVE  
L32      0 COUPLING (3A) MICROWAVE

|                      |                  |               |
|----------------------|------------------|---------------|
| COST IN U.S. DOLLARS | SINCE FILE ENTRY | TOTAL SESSION |
| FULL ESTIMATED COST  | 0.30             | 171.09        |

DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)

|                     |                  |               |
|---------------------|------------------|---------------|
| CA SUBSCRIBER PRICE | SINCE FILE ENTRY | TOTAL SESSION |
| 0.00                | -10.22           |               |

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FILE 'MEDLINE' ENTERED AT 12:33:10 ON 12 NOV 2005

FILE 'CAPLUS' ENTERED AT 12:33:10 ON 12 NOV 2005  
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=> coupling (3a) microwave  
L33      1138 COUPLING (3A) MICROWAVE

=> 133 and ("solid phase peptide" or SPPS)  
L34      6 L33 AND ("SOLID PHASE PEPTIDE" OR SPPS)

=> 134 and microwave

L35

6 L34 AND MICROWAVE

=> dup rem 135

PROCESSING COMPLETED FOR L35

L36 6 DUP REM L35 (0 DUPLICATES REMOVED)

=> d ibib abs total

L36 ANSWER 1 OF 6 CAPLUS COPYRIGHT 2005 ACS on STN

ACCESSION NUMBER: 2005:584968 CAPLUS

TITLE: Solid-phase synthesis of vapreotide via  
microwave

AUTHOR(S): Zhu, Yi-shen; Qiu, Qian; Tu, Chun-yan; Wei, Ping

CORPORATE SOURCE: College of Life Science and Pharmaceutical  
Engineering, Nanjing University of Technology,  
Nanjing, 210009, Peop. Rep. China

SOURCE: Jingxi Huagong (2005), 22(5), 395-397

CODEN: JIHUFJ; ISSN: 1003-5214

PUBLISHER: Jingxi Huagong Bianjibu

DOCUMENT TYPE: Journal

LANGUAGE: Chinese

AB Vapreotide was prepared by **solid-phase peptide**  
**synthesis** and effect of **microwave** on the **coupling**  
reaction was investigated by orthogonal test. Results were analyzed by  
multiple nonlinear regression method and response surface optimization.  
The optimal coupling reaction conditions were: maximum reaction temperature 60  
°C and reaction time 5 min, including 1 min for raising the temperature  
and 4 min for maintaining. Compared with the conventional method, the  
**microwave-enhanced coupling** reaction time was shortened  
about 12 .apprx. 36 times and less amts. of protected amino acids were  
needed. The yield of vapreotide was increased from 48% to 76%. It was  
identified by <sup>1</sup>H NMR, IR, MS and HRMS, and the results were consistent with  
the proposed structure.

L36 ANSWER 2 OF 6 CAPLUS COPYRIGHT 2005 ACS on STN

ACCESSION NUMBER: 2004:227419 CAPLUS

TITLE: Effect of **microwave** energy on **solid**  
**phase peptide** synthesis

AUTHOR(S): Collins, Jonathan M.; Hassman, C. Fred; King, Edward  
E.; Lambert, Joseph

CORPORATE SOURCE: Life Sciences Division, CEM Corporation, Matthews, NC,  
28106, USA

SOURCE: Abstracts of Papers, 227th ACS National Meeting,  
Anaheim, CA, United States, March 28-April 1, 2004  
(2004), ORGN-549. American Chemical Society:  
Washington, D. C.

CODEN: 69FGKM

DOCUMENT TYPE: Conference; Meeting Abstract

LANGUAGE: English

AB The application of **microwave** energy has proved to be a major  
enabling tool for many chemical applications requiring energy input. A new  
automated system for **microwave** assisted **solid**  
**phase peptide** synthesis has been developed that allows  
for complete cycle times of ten minutes as well as final peptide cleavage  
in ten minutes. A single mode cavity is used to allow for a high  
**microwave** power d. and a uniform field distribution. The  
stability of activated amino acids under **microwave** irradiation was  
investigated using PyBOP and HBTU activation. The effect of  
**microwave** energy on conventional side reactions with **SPPS**  
such as racemization and aspartimide formation was investigated and found  
to compare very favorably with conventional methods. Also, exciting  
changes in **coupling** chemistries possible with **microwave**  
energy will be presented that help to further suppress racemization. The  
application of this new method will be shown on a variety of peptide  
sequences.

L36 ANSWER 3 OF 6 CAPLUS COPYRIGHT 2005 ACS on STN  
ACCESSION NUMBER: 2004:658783 CAPLUS  
TITLE: **Microwave-enhanced solid-phase peptide synthesis**  
AUTHOR(S): Collins, Jonathan M.  
CORPORATE SOURCE: CEM Corporation, Matthews, NC, 28106-0200, USA  
SOURCE: Abstracts of Papers, 228th ACS National Meeting, Philadelphia, PA, United States, August 22-26, 2004 (2004), ORGN-518. American Chemical Society: Washington, D. C.  
CODEN: 69FTZ8  
DOCUMENT TYPE: Conference; Meeting Abstract  
LANGUAGE: English  
AB **Microwave** energy has proven to be a valuable tool for organic synthesis. Recently, **microwave** has been used for enhanced Fmoc **solid phase peptide** synthesis. With **microwave** energy, deprotection and **coupling** reactions can be performed in 3 and 4 min resp. This paper builds on previous work and demonstrates the successful application of **microwave** energy for longer 30-40 amino acid peptide sequences. Variation in deprotection and coupling chemistries will be presented and discussed.

L36 ANSWER 4 OF 6 CAPLUS COPYRIGHT 2005 ACS on STN  
ACCESSION NUMBER: 2002:674277 CAPLUS  
DOCUMENT NUMBER: 138:14167  
TITLE: **Rapid microwave-assisted solid phase peptide synthesis**  
AUTHOR(S): Erdelyi, Mate; Gogoll, Adolf  
CORPORATE SOURCE: Department of Organic Chemistry, Department of Medicinal Chemistry, Uppsala University, Uppsala, 751 21, Swed.  
SOURCE: Synthesis (2002), (11), 1592-1596  
CODEN: SYNTBF; ISSN: 0039-7881  
PUBLISHER: Georg Thieme Verlag  
DOCUMENT TYPE: Journal  
LANGUAGE: English  
OTHER SOURCE(S): CASREACT 138:14167  
AB A **microwave-assisted**, rapid **solid phase peptide** synthesis procedure is presented. It has been applied to the coupling of sterically hindered Fmoc-protected amino acids yielding di- and tripeptides. Optimized conditions for a variety of coupling reagents are reported. Peptides were obtained rapidly (1.5-20 min) and without racemization.  
REFERENCE COUNT: 19 THERE ARE 19 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L36 ANSWER 5 OF 6 CAPLUS COPYRIGHT 2005 ACS on STN  
ACCESSION NUMBER: 2000:294881 CAPLUS  
TITLE: **Coupling of near-grazing microwave photons to surface plasmon polaritons via a dielectric grating**  
AUTHOR(S): Hibbins, A. P.; Sambles, J. R.; Lawrence, C. R.  
CORPORATE SOURCE: School of Physics, Thin Film Photonics Group, University of Exeter, Exeter, EX4 4QL, UK  
SOURCE: Physical Review E: Statistical Physics, Plasmas, Fluids, and Related Interdisciplinary Topics (2000), 61(5-B), 5900-5906  
CODEN: PLEEE8; ISSN: 1063-651X  
PUBLISHER: American Physical Society  
DOCUMENT TYPE: Journal  
LANGUAGE: English  
AB A dielec. grating on top of a planar metal substrate is shown to couple near-grazing **microwave** photons to surface plasmon polaritons (**SPPs**). It is shown that when the grating grooves are oriented

such that they are parallel to the plane of incidence ( $\Phi=90^\circ$ ), coupling to SPPs with both s- and p-polarized photons is possible at three different energies. It is demonstrated that one mode is coupled via p-polarized radiation and the other two modes are both coupled via s-polarized radiation. A multilayer, multishape differential grating theory allows the identities of each of the modes to be confirmed by modeling the electromagnetic fields above the metal substrate. In addition, a comparison between the exptl. derived reflectivity scans and the theor. model is made.

REFERENCE COUNT: 18 THERE ARE 18 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L36 ANSWER 6 OF 6 CAPLUS COPYRIGHT 2005 ACS on STN

ACCESSION NUMBER: 1992:572007 CAPLUS

DOCUMENT NUMBER: 117:172007

TITLE: Enhanced coupling efficiency in solid-phase peptide synthesis by microwave irradiation

AUTHOR(S): Yu, Hui Ming; Chen, Shui Tein; Wang, Kung Tsung

CORPORATE SOURCE: Inst. Biol. Chem., Acad. Sin., Taipei, 10098, Taiwan

SOURCE: Journal of Organic Chemistry (1992), 57(18), 4781-4

CODEN: JOCEAH; ISSN: 0022-3263

DOCUMENT TYPE: Journal

LANGUAGE: English

AB Procedures have been developed for increasing coupling efficiency in solid-phase peptide synthesis by microwave irradiation using a kitchen microwave oven. A rate increase of at least 2-4 fold was observed. For side-chain hindered amino acids or for peptides containing difficult-coupling sequences, the peptide bond formation can be finished within 4-6 min. Under the same irradiation conditions, the microwave induced rate enhancement is more significant using Fmoc-peptide fragments than using amino acid derivs. in peptide synthesis. No detectable racemization reaction was observed

=> logoff h

COST IN U.S. DOLLARS

SINCE FILE

TOTAL  
ENTRY SESSION

FULL ESTIMATED COST

32.13

203.22

DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)

SINCE FILE

TOTAL  
ENTRY SESSION

CA SUBSCRIBER PRICE

-4.38

-14.60

SESSION WILL BE HELD FOR 60 MINUTES

STN INTERNATIONAL SESSION SUSPENDED AT 12:35:24 ON 12 NOV 2005